

B16

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

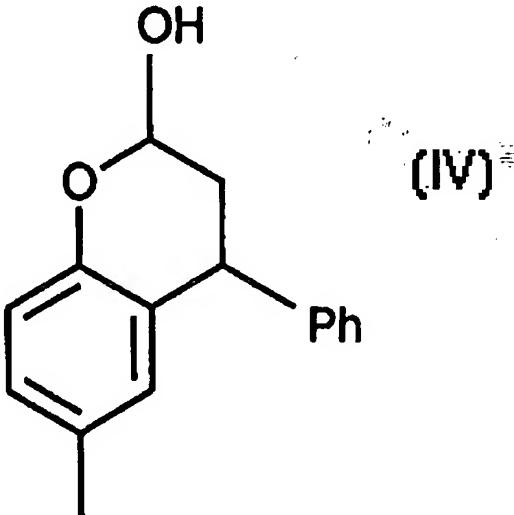
INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

| | | |
|---|---|--|
| (51) International Patent Classification ⁶ : | A1 | (11) International Publication Number: WO 98/29402 |
| C07D 311/20, C07C 215/54 | | (43) International Publication Date: 9 July 1998 (09.07.98) |
| (21) International Application Number: | PCT/US97/22521 | (81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). |
| (22) International Filing Date: | 18 December 1997 (18.12.97) | |
| (30) Priority Data: | 60/033,961 31 December 1996 (31.12.96) US | |
| (71) Applicant (for all designated States except US): | PHARMACIA & UPJOHN COMPANY [US/US]; 301 Henrietta Street, Kalamazoo, MI 49001 (US). | |
| (72) Inventors; and | | Published |
| (75) Inventors/Applicants (for US only): | GAGE, James, R. [US/US]; 341 Point-O-Woods Drive, Portage, MI 49002 (US). CABAJ, John, E. [US/US]; 4221 Autumn Court D-204, Sheboygan, WI 53081 (US). | With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments. |
| (74) Agent: | STEIN, Bruce; Pharmacia & Upjohn Company, Intellectual Property Legal Services, 301 Henrietta Street, Kalamazoo, MI 49001 (US). | |

(54) Title: PROCESS TO PREPARE TOLTERODINE

(57) Abstract

Disclosed is a novel intermediate, 3,4-dihydro-6-methyl-4-phenyl-2H-benzopyran-2-ol of formula (IV) and an improved process for the preparation of tolterodine.



FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

| | | | | | | | |
|----|--------------------------|----|---------------------------------------|----|---|----|--------------------------|
| AL | Albania | ES | Spain | LS | Lesotho | SI | Slovenia |
| AM | Armenia | FI | Finland | LT | Lithuania | SK | Slovakia |
| AT | Austria | FR | France | LU | Luxembourg | SN | Senegal |
| AU | Australia | GA | Gabon | LV | Latvia | SZ | Swaziland |
| AZ | Azerbaijan | GB | United Kingdom | MC | Monaco | TD | Chad |
| BA | Bosnia and Herzegovina | GE | Georgia | MD | Republic of Moldova | TG | Togo |
| BB | Barbados | GH | Ghana | MG | Madagascar | TJ | Tajikistan |
| BE | Belgium | GN | Guinea | MK | The former Yugoslav Republic of Macedonia | TM | Turkmenistan |
| BF | Burkina Faso | GR | Greece | | | TR | Turkey |
| BG | Bulgaria | HU | Hungary | ML | Mali | TT | Trinidad and Tobago |
| BJ | Benin | IE | Ireland | MN | Mongolia | UA | Ukraine |
| BR | Brazil | IL | Israel | MR | Mauritania | UG | Uganda |
| BY | Belarus | IS | Iceland | MW | Malawi | US | United States of America |
| CA | Canada | IT | Italy | MX | Mexico | UZ | Uzbekistan |
| CF | Central African Republic | JP | Japan | NE | Niger | VN | Viet Nam |
| CG | Congo | KE | Kenya | NL | Netherlands | YU | Yugoslavia |
| CH | Switzerland | KG | Kyrgyzstan | NO | Norway | ZW | Zimbabwe |
| CI | Côte d'Ivoire | KP | Democratic People's Republic of Korea | NZ | New Zealand | | |
| CM | Cameroon | | | PL | Poland | | |
| CN | China | KR | Republic of Korea | PT | Portugal | | |
| CU | Cuba | KZ | Kazakhstan | RO | Romania | | |
| CZ | Czech Republic | LC | Saint Lucia | RU | Russian Federation | | |
| DE | Germany | LI | Liechtenstein | SD | Sudan | | |
| DK | Denmark | LK | Sri Lanka | SE | Sweden | | |
| EE | Estonia | LR | Liberia | SG | Singapore | | |

WO 98/29402

PCT/US97/22521

PROCESS TO PREPARE TOLTERODINE

BACKGROUND OF THE INVENTION

5

1. Field of the Invention

The invention is a process to prepare tolterodine (V) including a novel intermediate. (R)-Tolterodine L-tartrate (VI) is useful in treating urinary incontinence.

2. Description of the Related Art

10 US 5,382,600 discloses tolterodine (V, and its tartrate salt) together with a method for its preparation. (R)-Tolterodine (VI) is useful for treating urinary incontinence.

Australian Journal of Chemistry, 26, 899-906 (1973) discloses the lactone (III).

SUMMARY OF INVENTION

15 Disclosed is 3,4-dihydro-6-methyl-4-phenyl-2H-benzopyran-2-ol (IV).

Also disclosed is a process for the production of tolterodine (V) which comprises:

(1) reducing the lactone (III) with a reducing agent to form the hydroxy compound (IV) and

20 (2) reductively aminating the hydroxy compound (IV) with diisopropylamine.

DETAILED DESCRIPTION OF THE INVENTION

US 5,382,600 discloses tolterodine (V, and its tartrate salt) together with a method for its preparation.

25 The improved process for preparing tolterodine is set forth in EXAMPLES 2 and 3.

Preferred reducing agents include diisobutylaluminum hydride, sodium bis(2-methoxyethoxy)aluminum and lithium tri-tert-butoxyaluminohydride; more preferred is diisobutylaluminum hydride (DIBAL).

It is preferred to perform the process of converting the lactone (III) to the corresponding hydroxy compound (IV) at temperatures of less than -15°; it is more preferable to perform this reaction at less than or equal to -20°.

30 The reaction of 4-hydroxytoluene (I) with the unsaturated acid (II) produces the lactone (III) which exists as two enantiomers. When the lactone (III) is reduced to the corresponding hydroxy compound (IV) the reduction of the carbonyl produces 35 a secondary alcohol with an stereogenic center. Hence, there are two pairs of diastereomers. Hence, when the term 3,4-dihydro-6-methyl-4-phenyl-2H-

WO 98/29402

PCT/US97/22521

benzopyran-2-ol (IV) is used it refers to and includes (2R,4R), (2S,4R), (2S,4S) and (2R,4S)-3,4-dihydro-6-methyl-4-phenyl-2H-benzopyran-2-ol (IV). In the transformation of the hydroxy compound (IV) to tolterodine (V), the center at 2 is lost producing tolterodine with one stereogenic center. This racemic compound is 5 later resolved in the conversion of tolterodine (V) to (R)-tolterodine L-tartrate (VI).

Tolterodine (V) is an amine, and as such forms acid addition salts when reacted with acids of sufficient strength. Pharmaceutically acceptable salts include salts of both inorganic and organic acids. The pharmaceutically acceptable salts are preferred over the corresponding free amines since they produce compounds which 10 are more water soluble and more crystalline. The preferred pharmaceutically acceptable salts include salts of the following acids methanesulfonic, hydrochloric, hydrobromic, sulfuric, phosphoric, nitric, benzoic, citric, tartaric, fumaric, maleic, $\text{CH}_3(\text{CH}_2)_n\text{-COOH}$ where n is 0 thru 4, $\text{HOOC}(\text{CH}_2)_n\text{-COOH}$ where n is as defined above. It is more preferred that the pharmaceutically acceptable salt of tolterodine 15 (V) is the tartrate (VI).

(R)-tolterodine L-tartrate (VI) is known to be useful for treating urinary incontinence.

DEFINITIONS AND CONVENTIONS

The definitions and explanations below are for the terms as used throughout 20 this entire document including both the specification and the claims.

I. CONVENTIONS FOR FORMULAS AND DEFINITIONS OF VARIABLES

The carbon atom content of variable substituents is indicated in one of two ways. The first method uses a prefix to the entire name of the variable such as " $\text{C}_1\text{-C}_4$ ", where both "1" and "4" are integers representing the minimum and maximum 25 number of carbon atoms in the variable. The prefix is separated from the variable by a space. For example, " $\text{C}_1\text{-C}_4$ alkyl" represents alkyl of 1 through 4 carbon atoms, (including isomeric forms thereof unless an express indication to the contrary is given). Whenever this single prefix is given, the prefix indicates the entire carbon atom content of the variable being defined. Thus $\text{C}_2\text{-C}_4$ alkoxy carbonyl describes a 30 group $\text{CH}_3(\text{CH}_2)_n\text{-O-CO-}$ where n is zero, one or two. By the second method the carbon atom content of only each portion of the definition is indicated separately by enclosing the " $\text{C}_i\text{-C}_j$ " designation in parentheses and placing it immediately (no intervening space) before the portion of the definition being defined. By this optional convention $(\text{C}_1\text{-C}_3)\text{alkoxycarbonyl}$ has the same meaning as $\text{C}_2\text{-C}_4$ alkoxy- 35 carbonyl because the " $\text{C}_1\text{-C}_3$ " refers only to the carbon atom content of the alkoxy group. Similarly while both $\text{C}_2\text{-C}_6$ alkoxyalkyl and $(\text{C}_1\text{-C}_3)\text{alkoxy}(\text{C}_1\text{-C}_3)\text{alkyl}$ define

WO 98/29402

PCT/US97/22521

alkoxyalkyl groups containing from 2 to 6 carbon atoms, the two definitions differ since the former definition allows either the alkoxy or alkyl portion alone to contain 4 or 5 carbon atoms while the latter definition limits either of these groups to 3 carbon atoms.

5

II. DEFINITIONS

All temperatures are in degrees Centigrade.

TLC refers to thin-layer chromatography.

HPLC refers to high pressure liquid chromatography.

Chromatography (column and flash chromatography) refers to

10 purification/separation of compounds expressed as (support, eluent). It is understood that the appropriate fractions are pooled and concentrated to give the desired compound(s).

$[\alpha]_D^{25}$ refers to the angle of rotation of plane polarized light (specific optical rotation) at 25° with the sodium D line (589A).

15 Pharmaceutically acceptable refers to those properties and/or substances which are acceptable to the patient from a pharmacological/toxicological point of view and to the manufacturing pharmaceutical chemist from a physical/chemical point of view regarding composition, formulation, stability, patient acceptance and bioavailability.

20 When solvent pairs are used, the ratios of solvents used are volume/volume (v/v).

DIBAL refers to diisobutylaluminum hydride.

EXAMPLES

Without further elaboration, it is believed that one skilled in the art can, 25 using the preceding description, practice the present invention to its fullest extent. The following detailed examples describe how to prepare the various compounds and/or perform the various processes of the invention and are to be construed as merely illustrative, and not limitations of the preceding disclosure in any way whatsoever. Those skilled in the art will promptly recognize appropriate variations 30 from the procedures both as to reactants and as to reaction conditions and techniques.

EXAMPLE 1 3,4-Dihydro-6-methyl-4-phenyl-2H-benzopyran-2-one (III)

Trans-cinammic acid (II, 100 g, 675 mmol) is added to a 1 L 4-neck round bottom flask equipped with a mechanical stirrer, thermocouple, and nitrogen inlet.

35 Para-cresol (I, 76.6 g, 708 mmol) is preheated in a water bath at 60° and added to the cinammic acid (II) followed by concentrated sulfuric acid (13.0 mL, 243 mmol).

WO 98/29402

PCT/US97/22521

The reaction is immediately heated to a set point of 122.5° and stirred at 120-125° until judged to be complete by HPLC analysis (column = nucleosil C-18; mobile phase = acetonitrile/water (45/55); flowrate = 1.5 ml/min; wavelength = 254 nm; sample preparation = (1) dissolve 6 drops of reaction mixture in methyl *t*-butylether (6 mL) pH 7 buffer, (2) dilute 0.4 mL of the organic layer in acetonitrile (5 mL) and inject;

5 retention times are: *t*-cinnamic acid = 3.3 min., *p*-cresol = 4.2 min. and the title compound = 20.3 min.) or TLC (acetone/cyclohexane (20/80), acetic acid (0.5%); wavelength = 254 nm) usually 6 hours. When the reaction is complete the mixture

10 is cooled to 100° and added to a prewarmed separatory funnel (500 mL). The bottom layer containing the sulfuric acid is removed and toluene (280 mL), water (50 mL) and potassium carbonate (47%, 10 mL) are added to the separatory funnel containing the crude product. If necessary the pH of the aqueous layer is adjusted to between 5-8 with additional 47% potassium carbonate. The layers are separated

15 and the organic layer is then washed once with water (50 mL). The organic layer is concentrated to a final volume of approximately 150 mL under reduced pressure. Isopropanol (350 mL) is then added, and distillation is continued to a volume of 350 mL. Isopropanol (150 mL) is again added and again distilled to 350 mL.

Isopropanol (150 mL) is again added and again distilled to 350 mL. The mixture is

20 then cooled to 30-40° with rapid stirring until the product crystallizes. The rapid stirring is continued after crystallization. The product is cooled to 0-5° and held at this temperature for approximately 1 hour, filtered and washed with isopropanol (200 mL) cooled to 0-5°. If the last portion of the wash is colored the wash is continued until no more color is removed. The solids are then dried at 60° under

25 reduced pressure to give the title compound, mp (uncorrected) = 83-85°.

EXAMPLE 2 3,4-Dihydro-6-methyl-4-phenyl-2H-benzopyran-2-ol (IV)

3,4-Dihydro-6-methyl-4-phenyl-2H-benzopyran-2-one (III, EXAMPLE 1, 100.0 g, 420.2 mmol) is added to toluene (500 mL). The mixture is degassed by purging alternately with vacuum and nitrogen and then cooled to -21°.

30 Diisobutylaluminumhydride in toluene solution (DIBAL, 1.5 M, 290 mL, 435 mmol) is then slowly added over 2 hr via add funnel while maintaining the reaction temperature at -20 to -25°. The reduction is usually done when the DIBAL add is completed. If the reaction is not done additional DIBAL can be added in increments. When the reaction is done (<1% lactone) ethyl acetate (45 mL) is added at -20° to -

35 25° via add funnel. Very little exotherm is observed. Next, citric acid (23%, 500 mL) is added. The mixture is stirred at 45-50° for 1 hr (or stirred overnight at 20-

WO 98/29402

PCT/US97/22521

- 25°), the phases are separated, the organic phase is washed with water (2 x 300 mL). The organic phase is concentrated to 250 mL under reduced pressure. Methanol (500 mL) is added, and the mixture is concentrated to 250 mL. The methanol addition and distillation is repeated to give the title compound in
- 5 methanol solution. This solution is concentrated to a thick oil which crystallizes on standing to give the title compound (as a mixture of diastereomers), IR (neat) 3410, 3020, 2925, 1605, 1498, 1447, 1205 and 1010 cm⁻¹; MS (m/z, EI) = 240 (parent). Rather than isolating and characterizing the title compound, it is normally taken directly into the next step.
- 10 HPLC (column = zorbax C-8; mobile phase acetonitrile/water (50/50); flow rate = 1 mL/min; wavelength = 240 nm; note - the absorbance of the lactone (III) at 240 nm is approximately 3.5 times greater than the lactol (IV); sample preparation is (1) add 3 drops of reaction mixture to methyl *t*-butyl ether (1 mL) and citric acid (23%, 1 mL) and shake for approximately 1 minute, (2) wash the organic phase once
- 15 with citric acid (23%, 1 mL) and once with water (1 mL), (3) dilute the organic phase (0.2 mL) in acetonitrile (1 mL) and inject; note - the methyl *t*-butyl ether layer must be sufficiently washed or an unknown peak at approximately 1.5 minutes will be present; retention times are: R_t (diol sideproduct) = 8.0 min, R_t (lactol II) = 15.9, 16.8 min (two diastereomers), R_t (lactone III) = 25.0 min.
- 20 EXAMPLE 3 Tolterodine hydrochloride (V)
- 3,4-Dihydro-6-methyl-4-phenyl-2H-benzopyran-2-ol (IV, EXAMPLE 2, 100 g) in methanol (500 mL) is slowly added to palladium on carbon (5%, 22 g, 10.5 mmol) while maintaining a slight nitrogen purge. If 3,4-dihydro-6-methyl-4-phenyl-2H-benzopyran-2-ol (IV) is added too quickly without a nitrogen purge the catalyst will
- 25 ignite the methanol. Diisopropylamine (147.0 mL, 1.05 mol) is added, and the mixture is hydrogenated at 45-50 psi and 48° until the reaction is judged to be complete by HPLC (< 2% lactol). The reaction is usually done after 10 hours, but can be run overnight. The reaction mixture is cooled and removed from the hydrogenator using a methanol (150 mL) rinse. The combined reaction mixture and
- 30 rinse is filtered through a bed of solka floc (10 g). The solka floc is washed thoroughly with methanol (100 mL) and the filtrate is concentrated to remove methanol while ethyl acetate is being added back. The volume of this solution of the free base of the title compound is adjusted to 700 mL using ethyl acetate and the mixture is heated to 55°.
- 35 To form the hydrochloride salt of the title compound, concentrated hydrochloric acid (52.5 mL, 630 mmol) is added over 15 min. The resulting slurry is

WO 98/29402

PCT/US97/22521

gradually cooled to -15° to -20° and held at this temperature for 1 hr. Tolterodine hydrochloride is collected by filtration, washed three times with ethyl acetate, and dried overnight under reduced pressure at 60° to give the title compound, mp = (uncorrected) 199-201°.

5 HPLC procedure is column = nucleosil C-18; mobile phase = acetonitrile/ammonium formate buffer (50/50) pH 3; flow rate = 1.5 mL/min; wavelength = 240 nm; retention times are R_t (tolterodine) = 8.7 min, R_t (diol sideproduct) = 7.3 min, R_t (lactol III) = 13.4 and 14.2 min (two diasteromers). Sample preparation is (1) dissolve 3 drops of the reaction mixture in methanol (1
10 mL), (2) filter through a syringe filter, (3) dilute filtered solution with acetonitrile (1 mL) and inject.

EXAMPLE 4 (R)-Tolterodine L-Tartrate (VI)

Tolterodine hydrochloride (V), EXAMPLE 3, 130.0 g, 359 mmol), methylene chloride (1.3 L) and water (650 mL) are mixed. The mixture is stirred rapidly while
15 adding sodium hydroxide (50%, 13.0 mL) and sodium carbonate (13.0 g, 123 mmol). The pH as determined by pH paper is 10-11. After stirring thoroughly for approximately 15 minutes two clear homogeneous phases form. Stirring is continued for another 45 minutes, the layers are separated and the organic phase is washed with water (2 x 650 mL). The methylene chloride mixture is concentrated
20 under reduced pressure. The concentrate is dissolved in ethanol (325 mL) and warmed to 60-70°. L-tartaric acid (80.84 g, 539 mmol) slurried in hot ethanol (810 mL) is added via add funnel at 60-70° over approximately 30 minutes. When the addition is done the slurry is refluxed for 1 hr, gradually cooled to 0° and held at this temperature for 1 hr. The slurry is filtered, washed with ethanol (2 x 260 mL)
25 previously cooled to 0°, and dried overnight under reduced pressure at 60° to give the crude title compound.

The crude product (136.0 g) and ethanol (5.44 L) are mixed and heated to 80° for 30 min. The mixture is concentrated to half the initial volume by distilling 2.72 L of ethanol. The mixture is gradually cooled to 20-25° over 1 hr, placed in an ice bath, and held at 0° for 1 additional hour. The tolterodine L-tartrate is collected by filtration, washed with ethanol (2 x 272 mL) previously cooled to 0°, and dried overnight under reduced pressure at 60° to give product. This procedure was repeated a second time on 81.0 g of once recrystallized tolterodine L-tartrate to give the optically active title compound, mp (uncorrected) = 210-211°; [α]²⁵ (1%,
35 methanol) = 27.4°.

EXAMPLE 5 Preparation of (R)-Tolterodine L-Tartrate In Methanol/Acetone

WO 98/29402

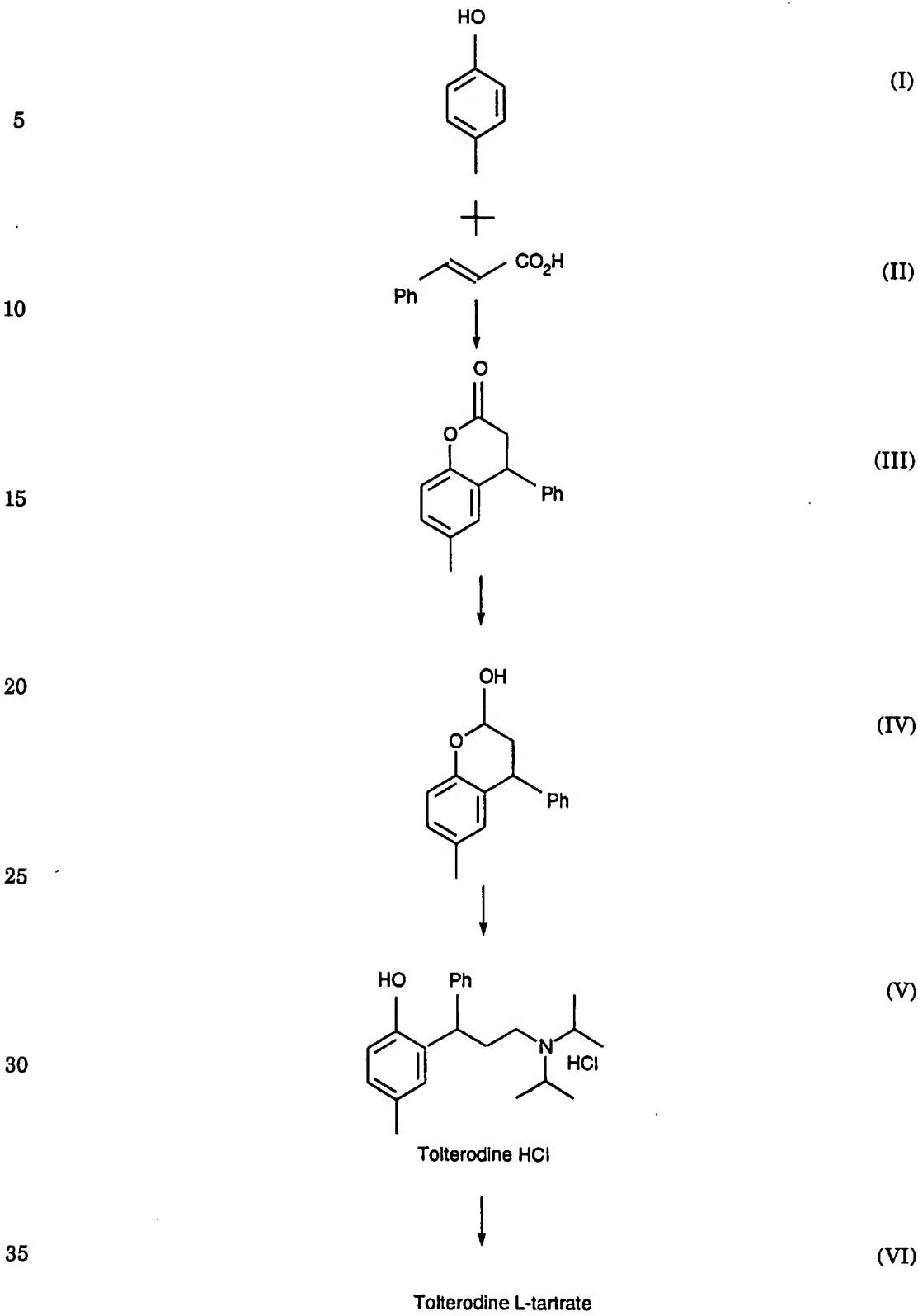
PCT/US97/22521

- Tolterodine hydrochloride (V), EXAMPLE 3, 130.0 g, 359 mmol), methylene chloride (1.3 L) and water (650 mL) are mixed. The thick slurry is stirred rapidly while adding sodium hydroxide (50%, 13.0 mL) and sodium carbonate (13.0 g, 123 mmol). The pH as determined by pH paper is 10-11. After stirring thoroughly for
5 approximately 15 minutes two clear homogeneous phases form. Stirring is continued for another 45 minutes, the layers are separated and the organic phase is washed with water (2 x 650 mL). The methylene chloride mixture is concentrated under reduced pressure. The concentrate is dissolved in acetone (1.3 L), warmed to 48-50° and L-tartaric acid (80.84 g, 539 mmol) slurried in hot methanol (162 mL) is
10 added via add funnel at 48-50° over approximately 30 minutes. The addition funnel is rinsed with acetone/methanol (90/10, 130 mL) and the slurry is refluxed for 1 hr before being gradually cooled to 0° for 1 hr. The mixture is filtered, washed with acetone (2 x 260 mL) previously cooled to 0°, and dried overnight in under reduced pressure at 60° to give crude (R)-tolterodine L-tartrate.
15 Crude (R)-tolterodine L-tartrate (115.0 g) and methanol (1.15 L) are slurried and heated to reflux for 30 min. The mixture is concentrated to half the initial volume by distilling 575 mL of methanol prior to adding acetone (3.26 L) over 30 min. The resulting slurry is refluxed for 1 hr and then gradually cooled to 20-25° over 1 hr before being placed in an ice bath and cooled to 0° for 1 additional hr. The
20 tolterodine L-tartrate is collected by filtration, washed with acetone (2 x 230 mL) previously cooled to 0°, and dried overnight under reduced pressure at 60° to give tolterodine L-tartrate. This procedure was repeated a second time on 82.0 g of once recrystallized tolterodine L-tartrate to give the optically active title compound, mp (uncorrected) = 210-211°; $[\alpha]^{25}$ (1%, methanol) = 27.4°.

WO 98/29402

PCT/US97/22521

CHART A



WO 98/29402

PCT/US97/22521

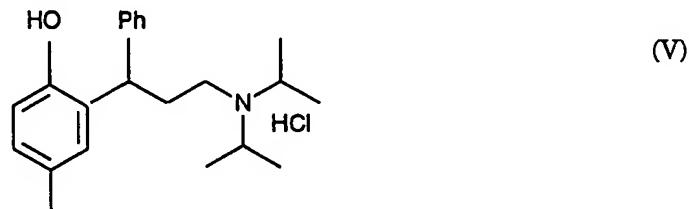
CLAIMS

1. 3,4-Dihydro-6-methyl-4-phenyl-2H-benzopyran-2-ol (IV).

2. A process for the production of tolterodine (V)

5

10



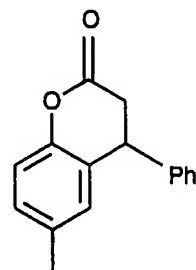
which comprises:

(1) reducing the lactone (III)

15

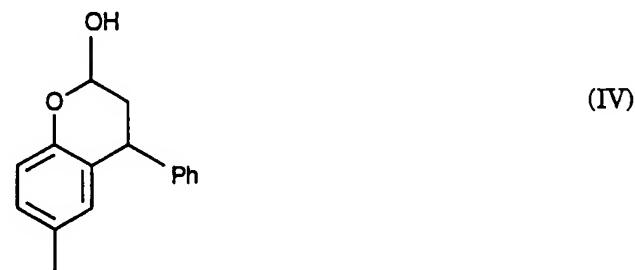
20

(III)



25 with a reducing agent to form the hydroxy compound (IV)

30



35 and

(2) reductively aminating the hydroxy compound (IV) with diisopropylamine.

WO 98/29402

PCT/US97/22521

3. A process according to claim 2 where the reducing agent is selected from the group consisting of diisobutylaluminum hydride, sodium bis(2-methoxyethoxy)aluminum and lithium tri-tert-butyloxyaluminohydride.
- 5 4. A process according to claim 2 where the reducing agent is diisobutylaluminum hydride.
5. A process according to claim 2 where step (1) is performed at less than -15°.
- 10 6. A process according to claim 5 where step (1) is performed at less than or equal to -20°.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 97/22521

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 C07D311/20 C07C215/54

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C07D C07C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

| Category | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
|----------|--|-----------------------|
| A | EP 0 325 571 A (KABIVITRUM) 26 July 1989 cited in the application see page 1 - page 8; claims ----- | 1, 2 |

 Further documents are listed in the continuation of box C. Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the International filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

13 May 1998

Date of mailing of the International search report

25/05/1998

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl.
Fax: (+31-70) 340-3016

Authorized officer

Francois, J

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 97/22521

| Patent document cited in search report | Publication date | Patent family member(s) | | Publication date |
|--|------------------|--|--|--|
| EP 325571 | A 26-07-1989 | AU AU DK EP HK HU HU JP JP NO WO US | 635493 B 2932989 A 172590 A 0354234 A 64494 A 212729 B 9400053 A 2664503 B 3503163 T 173496 C 8906644 A 5382600 A | 25-03-1993 11-08-1989 19-07-1990 14-02-1990 15-07-1994 28-10-1996 30-01-1995 15-10-1997 18-07-1991 22-12-1993 27-07-1989 17-01-1995 |